WEST Search History

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DATE: Thursday, July 06, 2006

Hide?	<u>Set</u> <u>Name</u>	Query	<u>Hit</u> Count
	DB=PG	SPB, USPT, DWPI; PLUR=NO; OP=ADJ	
	L19	(113 same antisense) and (@pd<20020210)	27
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	L17	L14 and ((pei or polyethyleneimine) with (encapsul\$ or cation\$ or lipos\$))	52 ·
	L16	L14 and ((pei or polyethyleneimine) with encapsul\$ or cation\$ or lipos\$)	3316
	L15	L14 and (pei or polyethyleneimine)	236
	L14	L13 and antisense	3555
	L13	((HA or NA or PB2 or PB1 or PA or NP or M1 or M2 or NS) with influenz\$)	5657
	L12	((PB2 or PB1 or PA or NP or M1 or M2 or NS) with influenz\$)	1333
	L11	110 and (pei near5 (cation\$ or lipos\$))	4
	L10	L9 and (@pd<20020210)	48
	L9	L8 and influenza .	396
	L8	antisense and pei	1750
	L7	l6 and (pei or polyethyleneimine)	11
	L6	15 and (cation\$ or lipid or (transfect near2 enhanc\$ or lipos\$))	51
[**]	L5	L4 and (sirna or shrna)	58
\Box .	L4	L3 and (influenz\$)	415
	L3	L2 and (influenz\$ or flu\$)	1094
	L2	L1 and (@pd<20030210 or @ad<20030210)	1170
	L1	(PTN or NP or influenz\$ or flu\$) and (sirna or rnai or dsrna or shrna)	3855

END OF SEARCH HISTORY

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SINCE FILE

ENTRY

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TOTAL

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SESSION

FULL ESTIMATED COST

COST IN U.S. DOLLARS

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FILE 'SCISEARCH' ENTERED AT 13:57:46 ON 06 JUL 2006 Copyright (c) 2006 The Thomson Corporation

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=> s l1 and l2

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Date specifications may use ranges and numeric operators. The date itself can be in any of the following general formats:

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Slash Format: MM/DD/YYYY or MM/YYYY

Dot Format: DD.MM.YYYY or MM.YYYY

Text Format: February 10, 1987 Feb 1989 Feb. 10, 1987 1990

Feb. 10, 2000 1998 - 2001

Feb 10, 1987 July 1997 - May 2002 10 February 1987 March 5 - 8, 1990 10 Feb 2007 April - June, 1999

Any year entered with only two digits will be interpreted as being in the range 1900-1999. Thus, Mar 12 01 will be searched as 19010312.

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=> d l19 bib abs 1-13

L19 ANSWER 1 OF 13 CA COPYRIGHT 2006 ACS on STN

AN 144:127482 CA

- TI Inhibition of class II antigen invariant chain (Ii) expression in mammalian cells using small interfering RNAs (siRNAs)
- IN Xu, Minzhen; Humphreys, Robert

PA USA

- SO U.S. Pat. Appl. Publ., 47 pp., Cont.-in-part of U.S. Ser. No. 127,347. CODEN: USXXCO
- DT Patent
- LA English

FAN.CNT 5

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	US 2002-54387	B2	20020122	•	
	US 2002-127347	A2	20020422		

- AB The present invention is directed toward compns. and methods involving the inhibition of Ii antigen (MHC Class II antigen invariant chain) expression in cells for the purpose of altering antigen presentation pathways. More specifically, disclosed are compns. and methods which relate to MHC Class II mol. presentation of antigenic epitopes which, under normal circumstances, would not be presented in association with MHC Class II mols. The invention relates to presentation in cells which normally express MHC Class II mols., as well as cells which can be induced to express MHC Class II mols. Adenoviral vectors are also constructed containing the transcription factor CIITA and/or interferon- γ cDNAs for tumor protection by MHC Class II+/Ii-. By suppressing the expression of the Ii protein, this vast repertoire of peptides which have been transported into the endoplasmic reticulum to MHC Class I and subsequent presentation to CD8+ T lymphocytes, can bind to MHC Class II mols. for subsequent presentation to, and activation of, CD4+ T immunoregulatory cells. Embodiments relating to RNA interference of Ii are specifically disclosed. Sequences of Ii-specific siRNAs are presented.
- L19 ANSWER 2 OF 13 CA COPYRIGHT 2006 ACS on STN
- AN 142:183225 CA
- TI RNA interference-mediated inhibition of gene expression using chemically modified short interfering nucleic acids
- IN McSwiggen, James; Chowrira, Bharat; Beigelman, Leonid; Macejak, Dennis; Zinnen, Shawn; Pavco, Pamela; Haeberli, Peter; Morissey, David; Fosnaugh, Kathy; Jamison, Sharon; Usman, Nassim; Thompson, James; Vargeese, Chandra; Wang, Weimen; Chen, Tonqian; Vaish, Narendra
- PA Sirna Therapeutics, Inc., USA
- SO U.S. Pat. Appl. Publ., 204 pp., Cont.-in-part of U.S. Ser. No. 720,448. CODEN: USXXCO
- DT Patent
- LA English

FAN.CNT 235

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AB The present invention concerns methods and reagents useful in modulating gene expression in a variety of applications, including use in therapeutic, diagnostic, target validation, and genomic discovery applications. Specifically, the invention relates to synthetic chemical modified small nucleic acid mols., such as short interfering nucleic acid (siNA), short interfering RNA (siRNA), double-stranded RNA (dsRNA), micro-RNA (miRNA), and short hairpin RNA (shRNA) mols. capable of mediating RNA interference (RNAi) against target nucleic acid sequences. Introduction of chemical modified nucleotides into nucleic acid mols. provides a powerful tool in overcoming potential limitations of in vivo stability and bioavailability inherent to native RNA mols. Unlike native unmodified siRNA, chemical modified siNA can also minimize the possibility of activating interferon activity in humans. Modifications are described including pyrimidine or purine nucleotides with 2'-deoxy-2'-fluoro or 2'-0-Me groups, phosphorothioate backbone modification, terminal residues comprising inverted deoxy thymidine or inverted deoxy abasic moieties, linking the sense and antisense strands with glyceryl succinate or dodecanoic acid or other linkers, and conjugation of targeting ligands (N-acetylgalactosamine, pteroic acid, peptides, or phospholipids) to the oligonucleotide termini. Thus, the serum stability of siNA constructs consisting of all RNA nucleotides containing two thymidine nucleotide overhangs have a half-life in human serum of 15 s, whereas chemical modified siNA constructs remained stable in serum for 1 to 3 days depending on the extent of modification. The small nucleic acid mols. are useful in the treatment of any disease or condition that responds to modulation of gene expression or activity in a cell, tissue, or organism. Three nuclease-resistant siNA mols. targeting site 1580 of hepatitis B virus RNA are designed using Stab 7/8 chemical and a 5'-terminal conjugate moiety (a branched cholesterol conjugate, a branched phospholipid conjugate, and a polyethylene glycol conjugate) showed significant stability in human and mouse serum (t1/2 = 10-408 h) and human liver extract (t1/2 = 28-43 h); the most stable siNA with all purine positions in the antisense strand with 2'-O-Me nucleotides had a half-life of 816 h in human liver extract

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TI RNA interference-mediated inhibition of gene expression using chemically modified short interfering nucleic acids

IN Mcswiggen, James; Chowrira, Bharat; Beigelman, Leonid; Macejak, Dennis; Zinnen, Shawn; Pavco, Pamela; Haeberli, Peter; Morrissey, David; Fosnaugh, Kathy; Jamison, Sharon; Usman, Nassim; Thompson, James; Vargeese, Chandra; Wang, Weimin; Chen, Tongqian; Vaish, Narendra

PA USA

SO U.S. Pat. Appl. Publ., 407 pp., Cont.-in-part of U.S. Ser. No. 427,160. CODEN: USXXCO

DT Patent

LA English

FAN.CNT 235

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The present invention concerns methods and reagents useful in modulating AΒ gene expression in a variety of applications, including use in therapeutic, diagnostic, target validation, and genomic discovery applications. Specifically, the invention relates to synthetic chemical modified small nucleic acid mols., such as short interfering nucleic acid (siNA), short interfering RNA (siRNA), double-stranded RNA (dsRNA), micro-RNA (miRNA), and short hairpin RNA (shRNA) mols. capable of mediating RNA interference (RNAi) against target nucleic acid Introduction of chemical modified nucleotides into nucleic acid mols. provides a powerful tool in overcoming potential limitations of in vivo stability and bioavailability inherent to native RNA mols. Unlike native unmodified siRNA, chemical modified siNA can also minimize the possibility of activating interferon activity in humans. Modifications are described including pyrimidine or purine nucleotides with 2'-deoxy-2'-fluoro or 2'-O-Me groups, phosphorothioate backbone modification, terminal residues comprising inverted deoxy thymidine or inverted deoxy abasic moieties, linking the sense and antisense strands with glyceryl succinate or dodecanoic acid or other linkers, and conjugation of targeting ligands (N-acetylgalactosamine, pteroic acid, peptides, or phospholipids) to the oligonucleotide termini. Thus, the serum stability of siNA constructs consisting of all RNA nucleotides containing two thymidine nucleotide overhangs have a half-life in human serum of 15 s, whereas chemical modified siNA constructs remained stable in serum for 1 to 3 days depending on the extent of modification. The small nucleic acid mols. are useful in the treatment of any disease or condition that responds to modulation of gene expression or activity in a cell, tissue, or organism. Three nuclease-resistant siNA mols. targeting site 1580 of hepatitis B virus RNA are designed using Stab 7/8 chemical and a 5'-terminal conjugate moiety (a branched cholesterol conjugate, a branched phospholipid conjugate, and a polyethylene glycol conjugate) showed significant stability in human and mouse serum (t1/2 = 10-408 h) and human liver extract (t1/2 = 28-43 h); the most stable siNA with all purine positions in the antisense strand with 2'-O-Me nucleotides had a half-life of 816 h in human liver extract

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TI RNA interference mediated inhibition of gene expression using chemically modified short interfering nucleic acid

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PA Ribozyme Pharmaceuticals, Incorporated, USA

SO PCT Int. Appl., 204 pp.
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AB The present invention concerns methods and reagents useful in modulating gene expression in a variety of applications, including use in therapeutic, diagnostic, target validation, and genomic discovery applications. Specifically, the invention relates to small nucleic acid mols., such as short interfering nucleic acid (siNA), short interfering RNA (siRNA), double-stranded RNA (dsRNA), micro-RNA (miRNA), and short hairpin RNA (shRNA) mols. capable of mediating RNA interference (RNAi) against target nucleic acid sequences. Exemplary siNA mols. are synthesized in tandem using standard phosphoramidite synthesis chemical

and a cleavable linker, for example a succinyl-based linker, followed by a one-step purification process that provides RNAi mols. in high yield. Chemical modifications (2'-O-Me and 2'-deoxy-2'-fluoro groups, phosphorothioate linkages, 5'-terminal caps comprising an inverted deoxy abasic moiety, etc.) in siNA constructs are selected to yield nuclease resistance while preserving the ability to mediate RNAi activity. The siNA mols. are designed that can bind to target vascular endothelial growth factor receptor R1 mRNA, hepatitis B virus RNA, and hepatitis C virus RNA, and are optionally individually analyzed by a computer folding algorithm to assess whether the siNA mol. can interact with the target sequence. The siNA mols. are useful in the treatment and diagnosis of prostate cancer and any other condition that responds to modulation of gene expression or activity in a cell, tissue, or organism.

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L19 ANSWER 5 OF 13 CA COPYRIGHT 2006 ACS on STN
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AN 138:249768 CA

TI Retroviral expression vectors for efficient transcription of small interfering RNAs in target cells

IN Baltimore, David; Qin, Xiao-Feng; Lois-Caballe, Carlos

PA California Institute of Technology, USA

SO PCT Int. Appl., 56 pp. CODEN: PIXXD2

DT Patent

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LA
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     WO 2003023015
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                        W
                               20020912
     WO 2002-US29214
                        W
                               20020913
     Retroviral expression vectors are described for use in the expression of
AB
     genes for small interfering RNAs in a target cell. Small interfering RNAs
     (siRNAs) that interfere with a viral life cycle by down
     regulating either the viral genome, a viral genome transcript, or an
     essential gene of a host cell may be transcribed from these vectors using
     a promoter for RNA polymerase III. In particular, these constructs can be
     used in the treatment of infection, particularly infection with HIV.
L19
    ANSWER 6 OF 13 CA COPYRIGHT 2006 ACS on STN
    138:142441 CA
AN
ΤI
    Enzymatic nucleic acid peptide conjugates
IN
    Beigelman, Leonid; Azhayev, Alex; Azhayeva, Elena
PA
    Ribozyme Pharmaceuticals, Inc., USA; Antopolsky, Maxim
SO
    PCT Int. Appl., 88 pp.
     CODEN: PIXXD2
DT
    Patent
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    English
FAN.CNT 1
                               DATE
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                                         APPLICATION NO.
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    WO 2003008628
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            FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR, BF, BJ, CF,
            CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
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US 2003148928

PRAI US 2001-306995P

A1

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US 2002-201389

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os MARPAT 138:142441

This invention features conjugates, compns., methods of synthesis, and AB applications thereof, including galactose, galactosamine, N-acetyl galactosamine, PEG, phospholipid, and human serum albumin (HSA) derived conjugates of nucleosides, nucleotides, non-nucleosides, and nucleic acids including enzymic nucleic acids, DNAzymes, allozymes, antisense, dsRNA, siRNA, triplex oligonucleotides, 2,5-A chimeras, decoys and aptamers.

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L19 ANSWER 7 OF 13 CA COPYRIGHT 2006 ACS on STN
AΝ
    138:131091 CA
ΤI
    Selective post-transcriptional silencing of oncogene in mammalian cells by
    siRNA for therapy
IN
    Milner, Anne Josephine
PΔ
    UK
SO
    PCT Int. Appl., 44 pp.
    CODEN: PIXXD2
DT
    Patent
    English
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FAN.CNT 1
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A WO 2002-GB3300 W 20020717 The present invention relates to a method of selective AB post-transcriptional silencing in a mammalian cell of the expression of an exogenous gene of viral origin. The method comprises introducing an siRNA construct into a mammalian cell where the siRNA construct is homologous to a part of the mRNA sequence of the exogenous

20020114

20020617

The invention also comprises an siRNA construct with a nucleotide sequence which is homologous to a part of the mRNA sequence of an exogenous gene of viral origin and to the use of such a construct as a medicament. SiRNA causes selective loss of HPV E6 and E7 mRNA.

E6 siRNA induces activation of P53 accompanied by p21

A

expression. E7 silencing results in de-phosphorylation of pRb and induces apoptosis of HPV-positives cells.

- L19 ANSWER 8 OF 13 MEDLINE on STN
- 2003106165 MEDLINE AΝ
- DN PubMed ID: 12594334

GB 2002-688

GB 2002-13855

- ΤI RNA interference of influenza virus production by directly targeting mRNA for degradation and indirectly inhibiting all viral RNA transcription.
- Ge Qing; McManus Michael T; Nguyen Tam; Shen Ching-Hung; Sharp ΑU

Phillip A; Eisen Herman N; Chen Jianzhu CS Center for Cancer Research and Department of Biology, Massachusetts Institute of Technology, 77 Massachusetts Avenue, Cambridge, MA 02139, USA. NC AI32486 (NIAID) AI40146 (NIAID) AI44477 (NIAID) AI44478 (NIAID) AI50631 (NIAID) CA42063 (NCI) CA60686 (NCI) GM34277 (NIGMS) SO Proceedings of the National Academy of Sciences of the United States of America, (2003 Mar 4) Vol. 100, No. 5, pp. 2718-23. Electronic Publication: 2003-02-19. Journal code: 7505876. ISSN: 0027-8424. CY United States DTJournal; Article; (JOURNAL ARTICLE) LA English FS Priority Journals EΜ 200305 ED Entered STN: 6 Mar 2003 Last Updated on STN: 14 May 2003 Entered Medline: 13 May 2003 AΒ Influenza A virus causes widespread infection in the human respiratory tract, but existing vaccines and drug therapy are of limited value. Here we show that short interfering RNAs (siRNAs) specific for conserved regions of the viral genome can potently inhibit influenza virus production in both cell lines and embryonated chicken eggs. The inhibition depends on the presence of a functional antisense strand in the siRNA duplex, suggesting that viral mRNA is the target of RNA interference. However, siRNA specific for nucleocapsid (NP) or a component of the RNA transcriptase (PA) abolished the accumulation of not only the corresponding mRNA but also virion RNA and its complementary RNA. These siRNAs also broadly inhibited the accumulation of other viral, but not cellular, RNAs. The findings reveal that newly synthesized NP and PA proteins are required for influenza virus transcription and replication and provide a basis for the development of sirNAs as prophylaxis and therapy for influenza infection in humans. ANSWER 9 OF 13 SCISEARCH COPYRIGHT (c) 2006 The Thomson Corporation on L19 AN2003:979106 SCISEARCH GA The Genuine Article (R) Number: 739WG Inhibition of virus replication by RNA interference ΤI ΑU Haasnoot J; Cupac D; Berkhout B (Reprint) CS Univ Amsterdam, Acad Med Ctr K3110, Dept Human Retrovirol, Meibergdreef 15, NL-1105 AZ Amsterdam, Netherlands (Reprint); Univ Amsterdam, Acad Med Ctr K3110, Dept Human Retrovirol, NL-1105 AZ Amsterdam, Netherlands CYA Netherlands SO JOURNAL OF BIOMEDICAL SCIENCE, (2003) Vol. 10, No. 6, Part 1, Sp. iss. SI, pp. 607-616. ISSN: 1021-7770. PR KARGER, ALLSCHWILERSTRASSE 10, CH-4009 BASEL, SWITZERLAND. General Review; Journal DT LA English REC Reference Count: 72 Entered STN: 21 Nov 2003 Last Updated on STN: 21 Nov 2003 *ABSTRACT IS AVAILABLE IN THE ALL AND IALL FORMATS* AΒ RNA interference (RNAi) is a sequence-specific gene-silencing mechanism in eukaryotes, which is believed to function as a defence

against viruses and transposons. Since its discovery, RNAi has been

developed into a widely used technique for generating genetic knock-outs and for studying gene function by reverse genetics. Additionally, inhibition of virus replication by means of induced RNAi has now been reported for numerous viruses, including several important human pathogens such as human immunodeficiency virus type 1, hepatitis C virus, hepatitis B virus, dengue virus, poliovirus and influenza virus A. In this review, we will summarize the current data on RNAi-mediated inhibition of virus replication and discuss the possibilities for the development of RNAi-based antiviral therapeutics. Copyright (C) 2003 National Science Council, ROC and S. Karger AG, Basel.

- L19 ANSWER 10 OF 13 MEDLINE on STN
- AN 2003416308 MEDLINE
- DN PubMed ID: 12954218
- TI The utility of siRNA transcripts produced by RNA polymerase i in down regulating viral gene expression and replication of negative- and positive-strand RNA viruses.
- AU McCown Matthew; Diamond Michael S; Pekosz Andrew
- CS Department of Molecular Microbiology, Washington University School of Medicine, St. Louis, MO 63110-1093, USA.
- SO Virology, (2003 Sep 1) Vol. 313, No. 2, pp. 514-24. Journal code: 0110674. ISSN: 0042-6822.
- CY United States
- DT Journal; Article; (JOURNAL ARTICLE)
- LA English
- FS Priority Journals
- EM 200310
- ED Entered STN: 5 Sep 2003 Last Updated on STN: 29 Oct 2003 Entered Medline: 28 Oct 2003
- AB Short interfering double-stranded RNAs (siRNAs) expressed under the control of an RNA polymerase I promoter system were used to target gene expression of influenza A and West Nile virus. Decreased RNA and protein expression was induced in a sequence-specific manner-reducing sequence complementarity from 21 to 17 nucleotides abrogated the siRNA effect. Reduced M(2) expression resulted in a decrease in total and infectious influenza A virus production. WNV protein expression, genomic RNA, and infectious virus production were all dramatically reduced by siRNAs targeting two distinct viral sequences. The data demonstrate the utility of plasmid-driven siRNAs in regulating the expression of single viral genes, global viral gene expression, as a potential antiviral treatment, and as a genetic tool for viruses whose genomes are difficult to manipulate.
- L19 ANSWER 11 OF 13 SCISEARCH COPYRIGHT (c) 2006 The Thomson Corporation on STN
- AN 2003:1044955 SCISEARCH
- GA The Genuine Article (R) Number: 739MR
- TI Improvement of the health status of livestock populations by reproductive-biotechnological technologies
- AU Niemann H (Reprint)
- CS FB Biotechnol, Inst Tierzucht, FAL Mariensee, D-31535 Neustadt, Germany (Reprint); FB Biotechnol, Inst Tierzucht, D-31535 Neustadt, Germany
- CYA Germany
- SO ZUCHTUNGSKUNDE, (SEP-OCT 2003) Vol. 75, No. 5, pp. 401-413. ISSN: 0044-5401.
- PB EUGEN ULMER GMBH CO, POSTFACH 700561 WOLLGRASWEG 41, D-70599 STUTTGART, GERMANY.
- DT Article; Journal
- LA German
- REC Reference Count: 42
- ED Entered STN: 9 Dec 2003
 - Last Updated on STN: 9 Dec 2003
 - *ABSTRACT IS AVAILABLE IN THE ALL AND IALL FORMATS*

AB Established reproductive-biotechnological technologies such as artificial insemination (AI) and embryo transfer (ET) employing in vivo produced embryos are already valuable tools for the maintenance and improvement of the health status of livestock populations. Transmission of pathogens can be avoided by employing strict hygienic protocols. The intact zona pellucida acts as a very efficient barrier against pathogenic infections in in vivo derived embryos. New hygienic challenges arise with the growing use of in vitro produced, cloned and/or transgenic embryos. But the extended experience gained from the global use of in vivo produced embryos forms a solid platform for eventual commercial applications of these types of embryos. However, as the zona pellucida is structurally different or not intact in these embryos major research is needed to ensure that no disease transmission occurs. Further options for the improvement of the health status of livestock populations will evolve from transgenic technologies, in particular when the genome of the major farm animals has been completely sequenced and mapped. Today there are already promising examples for the potential usefulness of this approach to generate disease resistant populations. In the future, gene therapy, DNA vaccination and small interfering RNAs (siRNAs) will also play an important role to maintain and improve the health of farm animals. is predicted that within the next 15-20 years a broad arsenal of technologies based on reproductive biological and molecular genetic tools will become available for improving animal health.

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L19 ANSWER 12 OF 13 MEDLINE on STN
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- AN 2003546501 MEDLINE
- DN PubMed ID: 14625885
- TI RNA interference: on the road to an alternate therapeutic strategy!.
- AU Dave Rajnish S; Pomerantz Roger J
- CS Center for Human Virology and Biodefense, Division of Infectious Diseases and Environmental Medicine, Thomas Jefferson University, Philadelphia, PA 19107, USA.
- NC AI43876 (NIAID)
 - MH58526 (NIMH)
 - NS27405 (NINDS)
 - NS44513 (NINDS)
- SO Reviews in medical virology, (2003 Nov-Dec) Vol. 13, No. 6, pp. 373-85. Ref: 96
 - Journal code: 9112448. ISSN: 1052-9276.
- CY England: United Kingdom
- DT Journal; Article; (JOURNAL ARTICLE)
 General Review; (REVIEW)
- LA English
- FS Priority Journals
- EM 200401
- ED Entered STN: 20 Nov 2003 Last Updated on STN: 8 Jan 2004 Entered Medline: 7 Jan 2004
- AB RNA interference (RNAi) is a newly described natural biological phenomenon mediated by small interfering RNA (siRNA) molecules which target viral mRNA for degradation by cellular enzymes. RNAi has become a method of choice for studying gene function, especially in mammalian systems. With proof-of-concept studies already presented against a wide variety of human pathogens and several innovative methods of delivering the siRNA to a wide variety of primary cells available, the role for siRNA as a potential therapeutic strategy is becoming increasingly clear. This review presents recent advances in this direction. Copyright 2003 John Wiley & Sons, Ltd.
- L19 ANSWER 13 OF 13 CA COPYRIGHT 2006 ACS on STN
- AN 138:14152 CA
- TI Preparation of enzymic ribonucleic acid peptide conjugates as antitumor and antiviral agents and compositions for cellular delivery
- IN Beigelman, Leonid; Matulic-Adamic, Jasenka; Vargeese, Chandra; Karpeisky,

Alexander; Blatt, Lawrence; Shaffer, Christopher Ribozyme Pharmaceuticals, Inc, USA PCT Int. Appl., 220 pp. CODEN: PIXXD2

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AB This invention features peptide nucleotide conjugates I wherein each R1-R8 are independently hydrogen, alkyl, substituted alkyl, aryl, substituted aryl, or a protecting group, each "n" is independently an integer from 0 to about 200, R9 is a straight or branched chain alkyl, substituted alkyl, aryl, or substituted aryl, and R2 is a phosphorus containing group, nucleoside, nucleotide, small mol., nucleic acid, or a solid support comprising a linker., degradable linkers, compns., methods of synthesis, and applications thereof, including folate, galactose, galactosamine, N-acetyl galactosamine, PEG, phospholipid, peptide and human serum albumin (HAS) derived conjugates of biol. active compds., including antibodies, antivirals, chemotherapeutics, peptides, proteins, hormones nucleosides, nucleotides, non-nucleosides, and nucleic acids including enzymic nucleic acids, DNAzymes, allozymes, antisense, dsRNA, siRNA, triplex oligonucleotides, 2,5-A chimeras, decoys and aptamers. Thus, 1-0-(4-monomethoxytrityl)-N-(12'-hydroxydodecanoyl-2-acetamido-3,4,6-tri-0acetyl-2-deoxy-3-D-galactopyranose) -D-threoninol 3-0-(2-cyanoethyl,N,Ndiisopropylphosphorami-dite) was prepared and incorporated into RNA. A method of treating a cancer patient, comprising contacting cells of patient wherein said cancer is breast cancer, lung cancer, colorectal cancer, brain cancer, esophageal cancer, stomach cancer, bladder cancer, pancreatic cancer, cervical cancer, head and neck cancer, ovarian cancer, melanoma, lymphoma, glioma, or multidrug resistant cancers and/or viral infections including HIV, HBV, HCV, CMV, RSV, HSV, poliovirus, influenza, rhinovirus, west nile virus, Ebola virus, foot and mouth virus, and papilloma.

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